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LETTER

COMPLEXO: identifying the missing heritability of breast cancer via next generation collaboration

COMPLEXO: Melissa C Southey^{1*}, Daniel J Park¹, Tu Nguyen-Dumont¹, Ian Campbell², Ella Thompson², Alison H Trainer³, Georgia Chenevix-Trench⁴, Jacques Simard⁵, Martine Dumont⁵, Penny Soucy⁵, Mads Thomassen⁶, Lars Jønson⁷, Inge S Pedersen⁸, Thomas VO Hansen⁷, Heli Nevanlinna⁹, Sofia Khan⁹, Olga Sinilnikova^{10,11}, Sylvie Mazoyer¹⁰, Fabienne Lesueur¹², Francesca Damiola¹⁰, Rita Schmutzler^{13,14}, Alfons Meindl¹⁵, Eric Hahnen^{13,14}, Michael R Dufault¹⁵, TL Chris Chan^{16,17}, Ava Kwong^{16,18}, Rosa Barkardóttir¹⁹, Paolo Radice²⁰, Paolo Peterlongo²¹, Peter Devilee²², Florentine Hilbers²², Javier Benitez²³, Anders Kvist²⁴, Therese Törngren²⁴, Douglas Easton²⁵, David Hunter²⁶, Sara Lindstrom²⁶, Peter Kraft²⁶, Wei Zheng²⁷, Yu-Tang Gao²⁸, Jirong Long²⁷, Susan Ramus²⁹, Bing-Jian Feng³⁰, Jeffrey N Weitzel³¹, Katherine Nathanson³², Kenneth Offit³³, Vijai Joseph³³, Mark Robson³³, Kasmintan Schrader³³, San Ming Wang³⁴, Yeong C Kim³⁴, Henry Lynch³⁵, Carrie Snyder³⁵, Sean Tavtigian³⁶, Susan Neuhausen³⁷, Fergus J Couch³⁸ and David E Goldgar³⁶

Linkage analysis, positional cloning, candidate gene mutation scanning and genome-wide association study approaches have all contributed significantly to our understanding of the underlying genetic architecture of breast cancer. Taken together, these approaches have identified genetic variation that explains approximately 30% of the overall familial risk of breast cancer, implying that more, and likely rarer, genetic susceptibility alleles remain to be discovered.

The application of massively parallel sequencing has further demonstrated the complexity of human genetic variation and has raised many challenges for computational and statistical methods for searching for additional breast cancer predisposition genes. Early findings are consistent with previous indications that no single gene is likely to account for a large proportion of the remaining unexplained genetic susceptibility [1,2].

Coordinated international collaboration offers great potential to advance the discovery of additional breast cancer susceptibility genes by increasing the likelihood of identifying functionally relevant genetic variants in the same genes in multiple families. A new consortium, COMPLEXO (a name chosen to reflect the complexity of the exome), has been formed to facilitate collaborations between researchers actively applying massively parallel

sequencing to understand the genetics of breast and ovarian cancer. The consortium has defined activities aimed at bringing together data and resources suitable for exome/genome sequencing initiatives and for large case-control-family study resources suitable for validation of candidate susceptibility genes in which rare mutations are associated with high to moderate risk of breast cancer. The aim of COMPLEXO is to bring to massively parallel sequencing the same power of large sample sets that have proven so successful in examining the role of common variants in cancer populations via the consortium model, such as the Breast Cancer Association Consortium (BCAC), the Consortium of Investigators of Modifiers of BRCA1/2 (CIMBA), the Ovarian Cancer Association Consortium (OCAC) and the Collaborative Oncology Gene-environment Study (COGS) [3-5]. However, sequencing studies provide additional challenges in terms of defining specific modes of collaboration given differences in sequencing and targeted capture platforms, bioinformatics platforms, the need to integrate ongoing studies in many centers and socio-ethical-legal issues that are not as relevant to initiatives that are genotyping common genetic variation.

COMPLEXO invites collaboration from any researcher who would like to contribute to this consortium either by contributing data to the combined COMPLEXO data set, contributing resources for large-scale validation of candidate breast cancer predisposition genes or refining analytical and bioinformatic pipelines for massively parallel sequencing data filtering and prioritization. COMPLEXO also has interests in the critical assessment of current platforms and protocols and in developing and improving

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data filtering and gene prioritization strategies to enhance gene discovery initiatives. These approaches are relevant to all complex human diseases.

Interested researchers can engage with COMPLEXO via any local member or by contacting the corresponding author.

Competing interests

The authors declare that they have no competing interests.

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References

1. Thompson ER, Doyle MA, Ryland GL, Rowley SM, Choong DY, Tothill RW, Thorne H; kConFab, Barnes DR, Li J, Ellul J, Philip GK, Antill YC, James PA, Trainer AH, Mitchell G, Campbell IG: **Exome sequencing identifies rare deleterious mutations in DNA repair genes FANCC and BLM as potential breast cancer susceptibility alleles.** *PLoS Genet* 2012, **8**:e1002894.
2. Park DJ, Lesueur F, Nguyen-Dumont T, Pertesi M, Odefrey F, Hammet F, Neuhausen SL, John EM, Andrulis IL, Terry MB, Daly M, Buys S, Le Calvez-Kelm F, Lonie A, Pope BJ, Tsimiklis H, Voegelé C, Hilbers FM, Hoogerbrugge N, Barroso A, Osorio A; Breast Cancer Family Registry; Kathleen Cunningham Foundation Consortium for Research into Familial Breast Cancer, Giles GG, Devilee P, Benitez J, Hopper JL, Tavtigian SV, Goldgar DE, Southey MC: **Rare mutations in XRCC2 increase the risk of breast cancer.** *Am J Hum Genet* 2012, **90**:734-739.
3. [http://ccge.medschl.cam.ac.uk/consortia/bcac/index.html]
4. **CIMBA (The Consortium of Investigators of Modifiers of BRCA1/2)** [http://ccge.medschl.cam.ac.uk/consortia/cimba/index.html]
5. **Collaborative Oncological Gene-environment Study** [http://www.cogseu.org]

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